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(54) Insecticidally active nitro compounds

Nitroverbindungen als Insektizide

Composés nitrés comme insecticide

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(Japan)

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CORRIGENDUM issued on 14.02.96

EP 0 375 907 B1

Description

The present invention relates to novel nitro compounds, to processes for their preparation and to their use as insecticides.

It has already been disclosed that a certain group of 2-nitro-1,1-ethenediamines is useful as medicaments which influence the circulation, in particular as hypotensive agents (see the U.S. Patent 4,567,188), a certain group of N-cyanoisothioureas is useful as medicaments for treating ulcers (see Japanese Patent Laid-open No. 234,064/1987), the N-cyanoisothioureas disclosed in the above Japanese patent application have also a function for controlling insects and plant-destructive nematodes (see Japanese Patent Laid-open No. 233,903 and EP-A 303,570), and that a certain group of α -unsaturated amines has insecticidal/miticidal activity (see EP-A 0302389).

The compounds 3-(3-pyridylmethyl)- and 3-(4-pyridylmethyl)-2-nitroguanidine have been published in Bulletin of National Institut of Hygienic Sciences (Japan) No. 88 (1970) and No. 102 (1989) without any indication as to their insecticidal properties.

There have now been found novel nitro compounds of the formula (I)



wherein R¹ and R² are hydrogen or C₁₋₄ alkyl,
R³ is -S-R⁴ or



in which R⁴ is

C₁₋₄ alkyl, R⁵ and R⁶ are hydrogen or C₁₋₄ alkyl,

Z is 3-pyridyl, 4-pyridyl or 5-thiazolyl which may be substituted by halogen or C₁₋₄-alkyl,

with the exception of 3-(3-pyridylmethyl)-2-nitroguanidine and 3-(4-pyridylmethyl)-2-nitroguanidine.

The compounds of the formula (I) can be obtained by a process in which

a): (in the case where R³ is -S-R⁴)
compounds of the formula (V)



wherein R² and R⁴ have the same meanings as mentioned above, are reacted with compounds of the formula (VI)



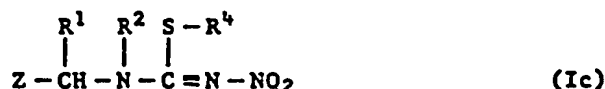
wherein R¹ and Z have the same meanings as mentioned above, and
Hal represents halogen atom,
in the presence of inert solvents, and if appropriate
in the presence of acid binders,

or

b): (in the case where R³ is



compounds of the formula (Ic)



wherein R¹, R², R⁴ and Z have the same meanings as mentioned above, are reacted with the compounds of formula (IV),



wherein R⁵ and R⁶ have the same meaning as above, in the presence of inert solvents.

The novel nitro compounds exhibit powerful insecticidal properties.

Surprisingly, the nitro compounds according to the invention exhibit a substantially greater insecticidal action than those known from the above-cited prior art.

In the formulae, the C₁₋₄ alkyl groups R¹, R², R⁴, R⁵ and R⁶ are straight-chain or branched, such as methyl, ethyl, n- and i-propyl and n-, i-, s- and t-butyl. Preferred is methyl.

In the formulae Z is particularly preferred as 3-pyridyl and 5-thiazolyl.

The substituent halogen atoms on Z are fluorine, chlorine, bromine and iodine. Preferred is chlorine.

The substituent C₁₋₄alkyl groups on Z are straight-chain or branched, such as methyl, ethyl, n- and i-propyl and n-, i-, s- and t-butyl. Preferred is methyl.

The substituent may be in all possible positions of Z.

Preferably it is in the 6 position of the 3-pyridyl-ring and in the 2 position of the 5-thiazolyl ring.

Among the nitro compounds according to the invention, of the formula (I), preferred compounds are those in which R¹ and R² are hydrogen or methyl,

R³ is -S-R⁴ or



in which R⁴ is methyl,

R⁵ and R⁶ are hydrogen or methyl,

and

Z is 3-pyridyl, 4-pyridyl or 5-thiazolyl which may be substituted by chlorine or methyl.

EP 0 375 907 B1

More preferred nitro compounds of the formula (I) are those in which R¹ and R² are hydrogen or methyl, R³ is



in which R⁵ and R⁶ are hydrogen or methyl, Z is 2-chloro-5-pyridyl or 2-chloro-5-thiazolyl.

Very particularly preferred nitro compounds of the formula (I) are those in which R¹ and R² are hydrogen or methyl, R³ is

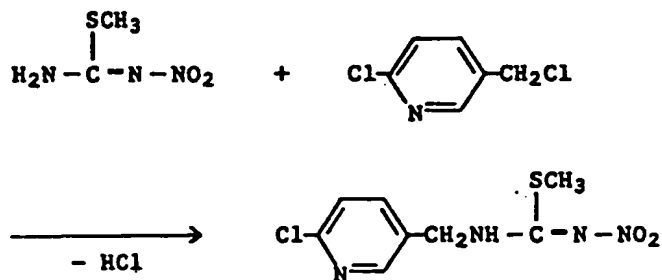


in which R⁵ and R⁶ are hydrogen or methyl, and Z is 2-chloro-5-pyridyl.

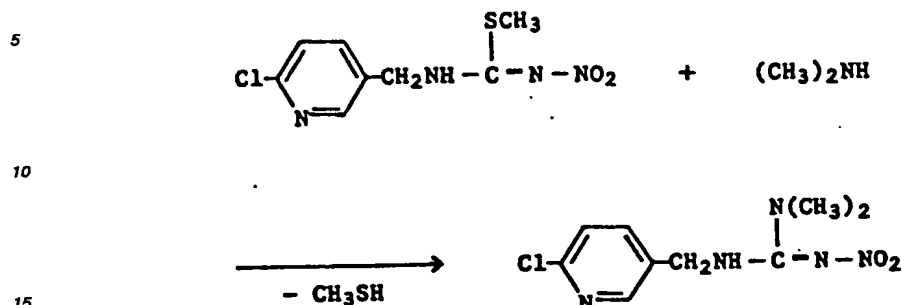
Specifically, the following compounds may be mentioned:

- 3-(2-chloro-5-pyridylmethyl)-1-methyl-2-nitroguanidine,
- 3-(2-chloro-5-pyridylmethyl)-1,1-dimethyl-2-nitroguanidine,
- 3-(2-chloro-5-pyridylmethyl)-1,3-dimethyl-2-nitroguanidine,
- 3-(2-chloro-5-pyridylmethyl)-1,1,3-trimethyl-2-nitroguanidine,
- 3-(2-chloro-5-pyridylmethyl)-3-methyl-2-nitroguanidine,
- 3-(2-chloro-5-thiazolylmethyl)-1-methyl-2-nitroguanidine.

If, for example, in the above process a), 2-methyl-3-nitroisothiourea and 2-chloro-5-chloromethylpyridine are used as starting materials, the course of the reaction can be represented by the following equation:



If, for example, in the above process b), 1-(2-chloro-5-pyridylmethyl)-3-nitro-2-methylisothiurea and dimethylamine are used as starting materials, the course of the reaction can be represented by the following equation:



In the process a), the compounds of the formula (V) as a starting material mean such ones based on the aforementioned definitions of R² and R⁴.

In the formula (V), R² and R⁴ preferably have the meanings already given above.

The compounds of the formula (V) are known (see e.g. J. Am. Chem. Soc., vol. 76, pages 1877 - 1879, 1954), and as examples, there may be mentioned: 3-nitro-2-methylisothiurea, 1,2-dimethyl-3-nitroisothiurea, 1,1,2-trimethyl-3-nitroisothiurea and so on.

The compounds of the formula (VI) as a starting material mean such ones based on the aforementioned definitions of R¹, Z and Hal.

In the formula (VI), R¹ and Z preferably have the meanings already given above, and Hal represents preferably chlorine or bromine.

The compounds of the formula (VI) are known (see Japanese Patent Laid-open Nos. 178981/1986, 178982/1986 or 183271/1986), and as examples, there may be mentioned: 2-chloro-5-chloromethylpyridine, 2-chloro-5-chloromethylthiazole and so on.

In the process b), the compounds of the formula (Ic) as a starting material correspond to a part of the compounds of the formula (I) which can be prepared by the above process a).

As compounds of the formula (IV) may be mentioned methylamine, diethylamine and so on.

Suitable diluents are all inert solvents.

These preferentially include water; aliphatic, cycloaliphatic and aromatic, optionally chlorinated, hydrocarbons, such as hexane, cyclohexane, petroleum ether, ligroin, benzene, toluene, xylene, methylene chloride, chloroform, carbon tetrachloride, ethylene chloride, trichloroethylene, chlorobenzene and the like; ethers such as diethyl ether, methyl ethyl ether, di-isopropyl ether, dibutyl ether, propylene oxide, dioxane, tetrahydrofuran and the like; ketones such as acetone, methyl-ethyl ketone, methyl-iso-propyl ketone, methyl-iso-butyl ketone; nitriles such as acetonitrile, propionitrile, acrylonitrile and the like; alcohols such as methanol, ethanol, iso-propanol, butanol, ethylene glycol and the like; esters such as ethyl acetate, amyl acetate; acid amides such as dimethyl formamide, dimethyl acetamide and the like; and sulfones and sulfoxides such as dimethyl sulfoxide, sulfolane and the like; and bases, for example, such as pyridine.

The above-mentioned process a) may be carried out in the presence of acid binders such as, for example, conventionally used hydroxide, hydride, carbonate, bicarbonate and alcoholate of alkali metal, tertiary amines such as, for example, triethyl amine, diethyl aniline, pyridine, etc.

In the above-mentioned process a), the reaction temperature can be varied within a wide range. In general, the reaction is carried out at a temperature of from about 0°C to boiling point of the reactant mixture, preferably from about 0°C to about 80°C.

In general, the reaction is preferably carried out under normal pressure, but can be verified also under elevated or reduced pressure.

In carrying out the process a), for instance, equi-mole to about 1.2 moles, preferably equi-mole of compound VI may be employed per mole of the compounds of the formula (V), and the mixture is reacted in the presence of inert solvents, such as e.g. dimethylsulfoxide and in the presence of acid binder, such as e.g. sodium hydride, so that the desired compounds of the formula (I) can be obtained.

In carrying out the process a), suitable diluents include the same solvents as exemplified for the process b).

In the above-mentioned process b), the reaction temperature can be varied within a wide range. In general, the reaction is carried out at a temperature in the range of from about 0°C to about 150°C, preferably from about 20°C to about 90°C.

In general, the reaction is preferably carried out under normal pressure, and also elevated or reduced pressure.

In carrying out the process b), for instance, equi-mole to slightly excessive mole amount of the compounds of the formula (IV) may be employed per mole of the compounds of the formula (Ic), and the mixture is reacted in the presence of inert solvents, so that the desired compounds of the formula (I) can be obtained.

The active compounds are well tolerated by plants, have a favorable level of toxicity to warm-blooded animals, and can be used for combating arthropod pests, especially insects which are encountered in agriculture, in forestry, in the protection of stored products and of materials, and in the hygiene field. They are active against normally sensitive and resistant species and against all or some stages of development. The above-mentioned pests include:

from the class of the Isopoda, for example Oniscus Asellus, Armadillidium vulgare and Porcellio scaber;

from the class of the Diplopoda, for example, Blaniulus guttulatus;

from the class of the Chilopoda, for example, Geophilus carpophagus and Scutigera spec.;

from the class of the Symphyla, for example, Scutigera immaculata;

from the order of the Thysanura, for example, Lepisma saccharina;

from the order of the Collembola, for example, Onychiurus armatus;

from the order of the Orthoptera; for example, Blatta orientalis, Periplaneta americana, Leucophaea maderae, Blattella germanica, Acheta domestica, Gryllotalpa spp., Locusta migratoria migratorioides, Melanoplus differentialis and Schistocerca gregaria;

from the order of the Dermaptera, for example, Forficula auricularia;

from the order of the Isoptera, for example, Reticulitermes spp.;

from the order of the Anoplura, for example, Phylloxera vastatrix, Pemphigus spp., Pediculus humanus corporis, Haematopinus spp. and Linognathus spp.;

from the order of the Mallophaga, for example, Trichodectes spp. and Damalinea spp.;

from the order of the Thysanoptera, for example, Hercinothrips femoralis and Thrips tabaci;

from the order of the Heteroptera, for example, Eurygaster spp., Dysdercus intermedius, Piesma quadrata, Cimex lectularius, Rhodnius prolixus and Triatoma spp.;

from the order of the Homoptera, for example, Aleurodes brassicae, Bemisia tabaci, Trialeurodes vaporariorum,

Aphis gossypii, Brevicoryne brassicae, Cryptomyzus ribis, Aphis fabae, Doralis pomi, Eriosoma lanigerum, Hyalopteris arundinis, Macrosiphum avenae, Myzus spp., Phorodon humuli, Rhopalosiphum padi, Empoasca spp., Euscelis bilobatus, Nephotettix cincticeps, Lecanium corni, Saundersia oleae, Laodelphax striatellus, Nilaparvata lugens, Aonidiella aurantii, Aspidiotus hederae, Pseudococcus spp. and Psylla spp.;

from the order of the Lepidoptera, for example, Pectinophora gossypiella, Bupalus piniarius, Cheimatobia brumata,

Lithocolletis blancardella, Hyponomeuta padella, Plutella maculipennis, Malcosoma neustria, Euproctis chrysorrhoea, Lymantria spp., Bucculatrix thurberiella, Phyllocnistis citrella, Agrotis spp., Euxoa spp., Feltia spp., Earias insulana, Heliothis spp., Spodoptera exigua, Mamestra brassicae, Panolis flammea, Prodenia litura, Spodoptera spp., Trichoplusia ni, Carpocapsa pomonella, Pieris spp., Chilo spp., Pyrausta nubilalis, Ephestia kuehniella, Galleria mellonella, Cacoecia podana, Capua reticulana, Choristoneura fumiferana, Clysia ambiguella, Homona magnanima and Tortrix viridana;

from the order of the Coleoptera, for example, Anobium punctatum, Rhizopertha dominica, Acanthoscelides obtectus, Acanthoscelides obtectus, Hylotrupes bajulus, Agelastica alni, Leptinotarsa decemlineata, Phaedon cochleariae, Diabrotica spp., Psylliodes chrysocephala, Epilachna varivestis, Atomaria spp., Oryzaephilus surinamensis, Anthonomus spp., Sitophilus spp., Otiorynchus sulcatus, Cosmopolites sordidus, Ceuthorrhynchus assimilis, Hypera postica, Dermestes spp., Trogoderma spp., Anthrenus spp., Attagenus spp., Lyctus spp., Meligethes aeneus, Ptinus spp., Niptus hololeucus, Gibbium psyllodes, Tribolium spp., Tenebrio molitor, Agriotes spp., Conoderus spp., Melolontha melolontha, Amphimallon solstitialis and Costelytra zealandica;

from the order of the Hymenoptera for example, Diprion spp., Hoplocamp spp., Lasius spp., Monomorium pharaonis and Vespa spp.;

from the order of the Diptera, for example, Aedes spp., Anopheles spp., Culex spp., Drosophila melanogaster, Musca spp., Fannia spp., Calliphora erythrocephala, Lucilia spp., Chrysomya spp., Cuterebra spp., Gastrophilus spp., Hypobosca spp., Stomoxys spp., Oestrus spp., Hypoderma spp., Tabanus spp., Tannia spp., Bibio hortulanus, Oscinella frit, Phorbia spp., Pegomya hyoscyami, Ceratitis capitata, Dacus oleae and Tipula paludosa;

from the order of the Siphonaptera, for example, Xenopsylla cheopis and Ceratophyllus spp.;

from the class of the Arachnida, for example, Scorpio maurus and Latrodectus mactans;

from the order of the Aranina, for example Acarus siro, Argas spp., Ornithodoros spp., Dermanyssus gallinae, Eriophyes ribis, Phyllocoptura oleivora, Boophilus spp., Rhipicephalus spp., Amblyomma spp., Hyalomma spp., Ixodes spp., Psoroptes spp., Chorioptes spp., Sarcoptes spp., Tarsonemus spp., Bryobia praetiosa, Panonychus spp. and Tetranychus spp..

The plant-parasitic nematodes include Pratylenchus spp., Radopholus similis, Ditylenchus dipsaci, Tylenchulus semipenetrans, Heterodera spp., Meloidogyne spp., Aphelenchoides spp., Longidorus spp., Xiphinema spp., and Trichodorus spp..

Furthermore, in the field of veterinary medicine, the novel compound of the present invention can effectively be employed for combating a variety of noxious animal-parasitic pests (internal- and external-parasitic pests), e.g., parasitic insects and nematodes. Such animal-parasitic pests may be exemplified as follows:

From the class of insects, e.g., Gastrophilus spp., Stomoxys spp., Tricodectes spp., Rhodius spp., Ctenocephalides canis and the like.

The active compounds can be converted into the customary formulations, such as solutions, emulsions, wettable powders, suspensions, powders, foams, pastes, granules, aerosols, natural and synthetic materials impregnated with active compound, very fine capsules in polymeric substances, coating compositions for use on seed, and formulations used with burning equipment, such as fumigating cartridges, fumigating cans and fumigating coils, as well as ULV cold mist and warm mist formulations.

These formulations may be produced in known manner, for example by mixing the active compounds with extenders, that is to say liquid or liquefied gaseous or solid diluents or carriers, optionally with the use of surface-active agents, that is to say emulsifying agents and/or dispersing agents and/or foam-forming agents. In the case of the use of water as an extender, organic solvents can, for example, also be used as auxiliary solvents.

As liquid solvents diluents or carriers, there are suitable in the main, aromatic hydrocarbons, such as xylene, toluene or alkyl naphthalenes, chlorinated aromatic or chlorinated aliphatic hydrocarbons, such as chlorobenzenes, chloroethylenes or methylene chloride, aliphatic hydrocarbons, such as cyclohexane or paraffins, for example mineral oil fractions, alcohols, such as butanol or glycol as well as their ethers and esters, ketones, such as acetone, methyl ethyl ketone, methyl isobutyl ketone or cyclohexanone, or strongly polar solvents, such as dimethylformamide and dimethylsulphoxide, as well as water.

By liquefied gaseous diluents or carriers are meant liquids which would be gaseous at normal temperature and under normal pressure, for example aerosol propellants, such as halogenated hydrocarbons as well as butane, propane, nitrogen and carbon dioxide.

As solid carriers there may be used ground natural minerals, such as kaolins, clays, talc, chalk, quartz, attapulgite, montmorillonite or diatomaceous earth, and ground synthetic minerals, such as highly-dispersed silicic acid, alumina and silicates. As solid carriers for granules there may be used crushed and fractionated natural rocks such as calcite, marble, pumice, sepiolite and dolomite, as well as synthetic granules of inorganic and organic meals, and granules of organic material such as sawdust, coconut shells, maize cobs and tobacco stalks.

As emulsifying and/or foam-forming agents there may be used non-ionic and anionic emulsifiers, such as polyoxyethylene-fatty acid esters, polyoxyethylene-fatty alcohol ethers, for example alkylaryl polyglycol ethers, alkyl sulphates, alkyl sulphates, aryl sulphonates as well as albumin hydrolysis products. Dispersing agents include, for example, lignin sulphite waste liquors and methylcellulose.

Adhesives such as carboxymethylcellulose and natural and synthetic polymers in the form of powders, granules or latices, such as gum arabic, polyvinyl alcohol and polyvinyl acetate, can be used in the formulation.

It is possible to use colorants such as inorganic pigments, for example iron oxide, titanium oxide and Prussian Blue, and organic dyestuffs, such as alizarin dyestuffs, azo dyestuffs or metal phthalocyanine dyestuffs, and trace nutrients, such as salts of iron, manganese boron, copper, cobalt, molybdenum and zinc.

The formulations in general contain from 0.1 to 95 per cent by weight of active compound, preferably from 0.5 to 90 per cent by weight.

The active compounds according to the invention can be present in their commercially available formulations and in the use forms, prepared from these formulations, as a mixture with other active compounds, such as insecticides, baits, sterilising agents, acaricides, nematicides, fungicides, growth-regulating substances or herbicides. The insecticides include, for example, phosphates, carbamates, carboxylates, chlorinated hydrocarbons, phenylureas, substances produced by microorganisms.

The active compounds according to the invention can furthermore be present in their commercially available formulations and in the use forms, prepared from these formulations, as a mixture with synergistic agents. Synergistic agent are compounds which increase the action of the active compounds, without it being necessary for the synergistic agent added to be active itself.

The active compound content of the use forms prepared from the commercially available formulations can vary within wide limits. The active compound concentration of the use forms can be from 0.0000001 to 100% by weight of active compound, preferably between 0.0001 and 1% by weight.

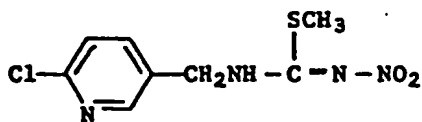
The compounds are employed in a customary manner appropriate for the use forms.

When used against hygiene pests and pests of stored products, the active compounds are distinguished by an excellent residual action on wood and clay as well as a good stability to alkali on limed substrates.

The preparation and use of the active compounds according to the invention can be seen from the following examples.

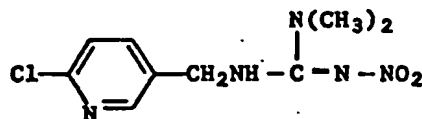
Examples

Example 1:



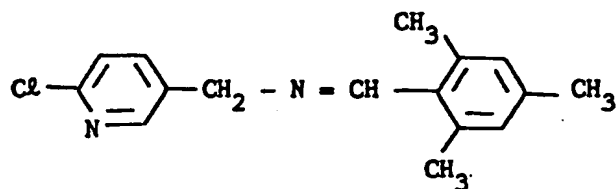
To a solution of 2-methyl-3-nitroisothiourea (15.0 g) in dimethylsulfoxide (100 ml) was gradually added sodium hydride (oil free 2.9 g) at 5°C, while being stirred for one hour. Thereafter, 2-chloro-5-chloromethyl pyridine (18.0 g) was added to the solution at a temperature in the range of from 5 to 10°C, followed by an overnight stirring thereof at room temperature. After the dimethylsulfoxide in the solution having been distilled off under reduced pressure, the resulting residue was purified on a chromatographic column (the eluent was a mixture of ethanol and chloroform), so as to obtain the desired 1-(2-chloro-5-pyridylmethyl)-2-methyl-3-nitroisothiourea (2.0 g) having a melting point in the range of from 141 to 143°C.

Example 2:



1-(2-chloro-5-pyridylmethyl)-2-methyl-3-nitroisothiourea (1.3 g) was dissolved in ethanol (20 ml) and to the solution was added an aqueous solution (50%) of dimethylamine (0.5 g) at room temperature, followed by a one-day stirring at 30°C. The ethanol in the solution was distilled off under reduced pressure and it was purified on a chromatographic column (the eluent was a mixture of methanol and chloroform) so as to obtain the desired 3-(2-chloro-5-pyridylmethyl)-1,1-dimethyl-2-nitroguanidine (1.2 g) having a melting point in the range of from 158 to 160°C.

Background example 1

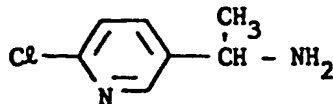


A mixture of 5-aminomethyl-2-chloropyridine (14.25 g), mesitaldehyde (14.8 g) and toluene (100 ml) was heated under reflux for 3 hours. The water which was formed during the reaction was trapped by Dean and Stark constant water separator.

The toluene was distilled off from the mixture under reduced pressure, so as to obtain N-(2,4,6-trimethylbenzylidene)-2-chloro-5-pyridylmethylaniline (27 g). m.p. 47 - 48°C.

Background example 2

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To a stirred solution of N-(2,4,6-trimethylbenzylidene)-2-chloro-5-pyridylmethanamine (10.9 g) in dried tetrahydrofuran (150 ml) cooled to -70°C under N₂ atmosphere was added butyl-lithium (10 w/v% in hexane, 26 ml), causing an immediate intense colour to occur. After 30 minutes stirring at the temperature, methyl iodide (5.7 g) was added dropwise, and then the mixture was stirred for three hours without cooling. The solvent was removed under reduced pressure.

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To the residue, ethanol (50 ml) and 2N-hydrochloric acid (50 ml) were added and heated under reflux for one hour.

The cooled solution was poured into water (100 ml) and extracted with methylene chloride (50 ml x 2).

The aqueous fraction was neutralized with 2N-sodium hydroxide and extracted with methylene chloride (50 ml x 2). The extract was dried over sodium sulfate and evaporated to obtain 1-(2-chloro-5-pyridyl)ethanamine (4 g).

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n_D^{20} 1.5440

The compounds of the formula (I), according to the invention which can be prepared in the same way as in

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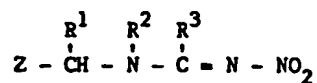
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Examples 1 to 2 are in the following Table 1.

Table 1



Compd. No.	Z	R ¹	R ²	R ³	
12	2-pyridyl	H	H	SCH ₃	
13	3-pyridyl	H	H	SCH ₃	
14	4-pyridyl	H	H	SCH ₃	
15	2-chloro-5-pyridyl	H	H	SCH ₃	m.p. 141 - 143°C
16	2-fluoro-5-pyridyl	CH ₃	H	SCH ₃	
17	2-chloro-5-pyridyl	H	CH ₃	SCH ₃	
18	2-methyl-5-pyridyl	H	H	SC ₂ H ₅	
19	2-chloro-5-pyridyl	CH ₃	CH ₃	SC ₂ H ₅	
20	2-bromo-5-pyridyl	H	H	SC ₃ H _{7-n}	
21	2-chloro-5-pyridyl	H	H	NH ₂	m.p. 197 - 199°C
22	2-chloro-5-pyridyl	H	CH ₃	NH ₂	m.p. 166 - 168°C
23	2-chloro-5-pyridyl	H	C ₂ H ₅	NH ₂	
24	2-chloro-5-pyridyl	H	C ₃ H _{7-n}	NH ₂	

EP 0 375 907 B1

Compd. No.	Z	R ¹	R ²	R ³	
25	2-chloro-5-pyridyl	CH ₃	H	NH ₂	n _D ²⁰ 1.5739
26	2-bromo-5-pyridyl	H	H	NH ₂	
27	2-chloro-5-thiazolyl	H	H	NH ₂	
28	2-chloro-5-pyridyl	H	H	NHCH ₃	m.p. 151 - 155°C
29	2-chloro-5-pyridyl	H	CH ₃	NHCH ₃	m.p. 135 - 139°C
30	2-chloro-5-pyridyl	H	C ₂ H ₅	NHCH ₃	
31	2-chloro-5-pyridyl	H	C ₃ H _{7-n}	NHCH ₃	
32	2-chloro-5-pyridyl	CH ₃	H	NHCH ₃	
34	2-chloro-5-thiazolyl	H	CH ₃	NHCH ₃	
35	2-chloro-5-pyridyl	H	H	NHC ₂ H ₅	m.p. 123 - 127°C
36	2-chloro-5-pyridyl	H	CH ₃	NHC ₂ H ₅	
37	2-chloro-5-pyridyl	C ₄ H _{9-n}	C ₂ H ₅	NHC ₂ H ₅	
38	2-fluoro-5-pyridyl	H	H	NHC ₂ H ₅	
39	2-chloro-5-pyridyl	H	H	NHC ₃ H _{7-n}	
40	2-chloro-5-pyridyl	H	CH ₃	NHC ₃ H _{7-n}	

EP 0 375 907 B1

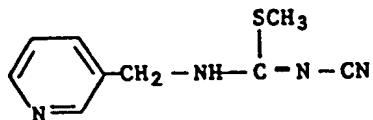
Compd. No.	Z	R ¹	R ²	R ³	
41	2-chloro-5-thiazolyl	H	H	NHC ₃ H ₇ -n	
42	2-chloro-5-pyridyl	H	C ₂ H ₅	NHC ₃ H ₇ -n	
43	2-chloro-5-pyridyl	H	H	NHC ₃ H ₇ -iso	m.p. 161 - 165°C
44	2-chloro-5-pyridyl	H	CH ₃	NHC ₃ H ₇ -iso	
45	2-chloro-5-pyridyl	H	H	N(CH ₃) ₂	m.p. 158 - 160°C
46	2-chloro-5-thiazolyl	H	H	N(CH ₃) ₂	
47	2-chloro-5-thiazolyl	H	CH ₃	N(CH ₃) ₂	
48	2-chloro-5-pyridyl	H	CH ₃	N(CH ₃) ₂	m.p. 96 - 99°C
49	2-chloro-5-pyridyl	H	C ₂ H ₅	N(CH ₃) ₂	
50	2-chloro-5-thiazolyl	CH ₃	H	N(CH ₃) ₂	
51	2-chloro-5-pyridyl	H	H	N-C ₂ H ₅ CH ₃	
52	2-chloro-5-pyridyl	H	CH ₃	N-C ₂ H ₅ CH ₃	
53	2-chloro-5-thiazolyl	H	H	N-C ₂ H ₅ CH ₃	
54	2-chloro-5-thiazolyl	H	CH ₃	N-C ₂ H ₅ CH ₃	

Compd. No.	Z	R ¹	R ²	R ³	
55	2-chloro-5-pyridyl	H	H	N-C ₃ H ₇ -iso CH ₃	
56	2-chloro-5-pyridyl	H	H	N(C ₂ H ₅) ₂	
57	2-chloro-5-pyridyl	H	CH ₃	N(C ₂ H ₅) ₂	
58	2-chloro-5-pyridyl	H	C ₂ H ₅	N(C ₂ H ₅) ₂	
59	2-chloro-5-pyridyl	H	C ₄ H ₉ -n	NHCH ₃	
60	2-chloro-5-pyridyl	H	H	NHC ₄ H ₉ -n	
61	4-pyridyl	H	CH ₃	NH ₂	m.p. 153 - 155°C
62	2-pyridyl	H	CH ₃	NH ₂	
63	3-pyridyl	H	CH ₃	NH ₂	m.p. 139 - 141°C
64	4-pyridyl	H	H	NHCH ₃	
65	4-pyridyl	H	CH ₃	NHCH ₃	
66	3-pyridyl	H	H	NHCH ₃	
67	3-pyridyl	H	CH ₃	NHCH ₃	
68	2-pyridyl	H	H	NHCH ₃	
69	2-pyridyl	H	CH ₃	NHCH ₃	

Compd. No.	Z	R ¹	R ²	R ³	
70	4-pyridyl	H	H	N(CH ₃) ₂	
71	3-pyridyl	H	H	N(CH ₃) ₂	
72	2-pyridyl	H	H	N(CH ₃) ₂	
73	2-methyl-5-pyrazinyl	H	CH ₃	NH ₂	
74	2-pyrimidinyl	H	H	NH ₂	
75	3-methyl-5-isoxazolyl	H	H	NH ₂	
76	3-methyl-5-isoxazolyl	H	H	NHCH ₃	
77	3-methyl-5-isoxazolyl	H	H	N(CH ₃) ₂	

Biological tests

Comparative compound E-1



(disclosed in Japanese Patent
Laid-open No. 233903/1988)

Example A (biological test)

Test on Nephotettix cincticeps having resistance to organophosphorus agents:-

Preparation of a test chemical

Solvent: 3 parts by weight of xylene
Emulsifier: 1 part by weight of polyoxyethylene alkyl phenyl ether

To form a suitable preparation, 1 part by weight of the active compound was mixed with the aforesaid amount of the solvent containing the aforesaid amount of the emulsifier. The mixture was diluted with water to a predetermined concentration.

Testing method

Onto rice plants, about 10 cm tall, planted in pots each having a diameter of 12 cm was sprayed 10 ml per pot of the water-dilution of each active compound in a predetermined concentration prepared as above. The sprayed chemical was dried, and a wire net having a diameter of 7 cm and a height of 14 cm was put over each pot, and 30 female imagoes of Nephotettix cincticeps showing resistance to organophosphorus agents were released into the net. The pots were each placed in a constant temperature chamber and the number of dead insects was examined 2 days later, and the insect mortality was calculated.

The results are shown in Table a.

Table a

Compound No.	Concentration of the active component, ppm	Insect mortality, %
22	8	100
45	8	100
Control E-1	50	0

Example B (biological test)

Test on planthoppers:-

5 Testing method

A water dilution in a predetermined concentration of the active compound prepared as in Example A was sprayed onto rice plants, about 10 cm tall, grown in pots with a diameter of 12 cm in an amount of 10 ml per pot. The sprayed chemical was dried, and a wire net, 7 cm in diameter and 14 cm tall, was put over each of the pots. Thirty female imagoes of *Nilaparvata lugens* Stal of a strain which showed resistance to organophosphorus chemicals were released into the net. The pots were left to stand in a constant temperature chamber and the number of dead insects was examined two days later. The kill ratio was then calculated.

In the same way as above, the insect mortality was calculated on *Sogatella furcifera* Horvath and organophosphorus-resistant *Laodelphax striatellus* Fallen.

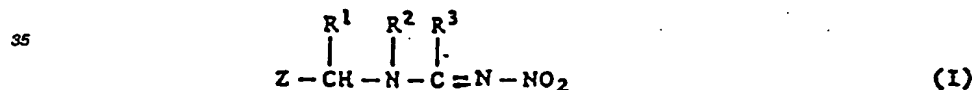
15 The results are shown in Table b.

Table b

Compound No.	Concentration of the active component, ppm	Insect mortality, %		
		<i>Nilaparvata lugens</i>	<i>Laodelphax striatellus</i>	<i>Sogatella furcifera</i>
22	8	100	100	100
45	8	100	100	100
Control E-1	50	0	0	0

30 Claims

1. Nitro compounds of the formula (I)



wherein R¹ and R² are hydrogen or C₁₋₄ alkyl,
R³ is -S-R⁴ or



in which R⁴ is C₁₋₄ alkyl, R⁵ and R⁶ are hydrogen or C₁₋₄ alkyl,
Z is 3-pyridyl, 4-pyridyl or 5-thiazolyl which may be substituted by halogen or C₁₋₄-alkyl,
with the exception of 3-(3-pyridylmethyl)-2-nitroguanidine and 3-(4-pyridylmethyl)-2-nitroguanidine.

2. Compounds of the formula (I) according to claim 1 wherein
R¹ and R² are hydrogen or methyl,

R³ is



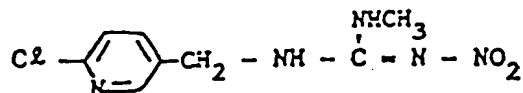
in which R⁵ and R⁶ are hydrogen or methyl,
Z is 2-chloro-5-pyridyl or 2-chloro-5-thiazolyl

3. Compounds of the formula (I) according to claim 1 wherein
R¹ and R² are hydrogen or methyl,
R³ is

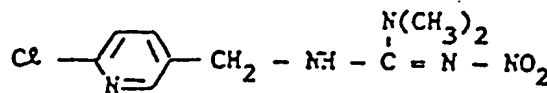


in which R⁵ and R⁶ are hydrogen or methyl,
Z is 2-chloro-5-pyridyl.

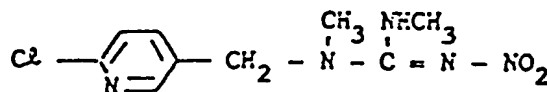
4. Compounds of the formula (I) according to claim 1 having the following individual structures:
3-(2-chloro-5-pyridylmethyl)-1-methyl-2-nitroguanidine represented by the following formula



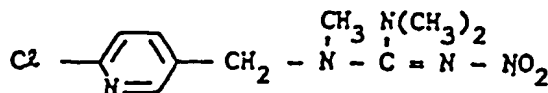
3-(2-chloro-5-pyridylmethyl)-1,1-dimethyl-2-nitroguanidine represented by the following formula



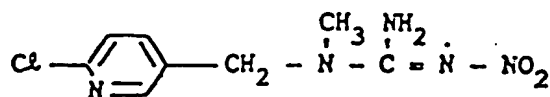
3-(2-chloro-5-pyridylmethyl)-1,3-dimethyl-2-nitroguanidine represented by the following formula



3-(2-chloro-5-pyridylmethyl)-1,1,3-trimethyl-2-nitroguanidine represented by the following formula



3-(2-chloro-5-pyridylmethyl)-3-methyl-2-nitroguanidine represented by the following formula



5. Process for the preparation of nitro compounds of the formula (I)



wherein R¹ and R² are hydrogen or C₁₋₄ alkyl,
R³ is -S-R⁴ or



in which R⁴ is
C₁₋₄ alkyl, R⁵ and R⁶ are hydrogen C₁₋₄ alkyl,
Z is 3-pyridyl, 4-pyridyl or 5-thiazolyl
which may be substituted by halogen or C₁₋₄-alkyl,
with the exception of 3-(3-pyridylmethyl)-2-nitroguanidine and 3-(4-pyridylmethyl)-2-nitroguanidine,

a) (in the case where R³ is -S-R⁴)
compounds of the formula (V)

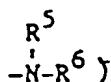


wherein R² and R⁴ have the same meanings as mentioned above, are reacted with compounds of the
formula (VI)

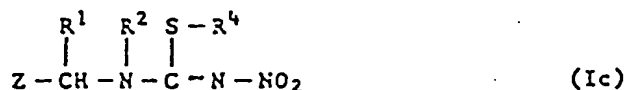


wherein R¹ and Z have the same meanings as mentioned above, and
Hal represents halogen atom,
in the presence of inert solvents, and if appropriate
in the presence of acid binders,

b) (in the case where R³ is



compounds of the formula (Ic)



wherein R¹, R², R⁴ and Z have the same meanings as mentioned above, are reacted with the compounds of formula (IV),

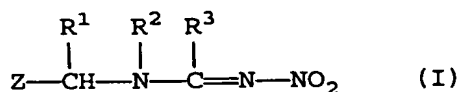


wherein R⁵ and R⁶ have the same meaning as above, in the presence of inert solvents.

6. Insecticidal compositions, characterised in that they contain at least one nitro compound of the formula (I), according to claims 1-4
7. Process for the preparation of insecticidal compositions, characterised in that nitro compounds of the formula (I), according to claims 1-4 are mixed with extenders and/or surface-active agents.
8. Method of combating harmful insects by applying to them or their habitat nitro compounds of the formula I according to claims 1-4.

Patentansprüche

1. Nitro-Verbindungen der Formel (I)



worin

R¹ und R² Wasserstoff oder C₁₋₄-Alkyl sind,
R³ -S-R⁴ oder



ist, worin
C₁₋₄-Alkyl ist
R⁴ Wasserstoff oder C₁₋₄-Alkyl sind,
R⁵ und R⁶ 3-Pyridyl, 4-Pyridyl oder 5-Thiazolyl ist, das durch Halogen oder C₁₋₄-Alkyl substituiert sein kann,
Z

mit Ausnahme von 3-(3-Pyridylmethyl)-2-nitroguanidin und 3-(4-Pyridylmethyl)-2-nitroguanidin.

2. Verbindungen der Formel (I) nach Anspruch 1, worin

R¹ und R² Wasserstoff oder Methyl sind,
R³



ist, worin

R⁵ und R⁶ Wasserstoff oder Methyl sind,
Z 2-Chlor-5-pyridyl oder 2-Chlor-5-thiazolyl ist.

3. Verbindungen der Formel (I) nach Anspruch 1, worin

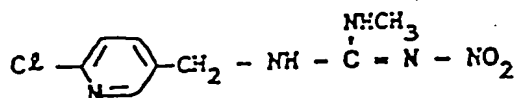
R¹ und R² Wasserstoff oder Methyl sind,
R³



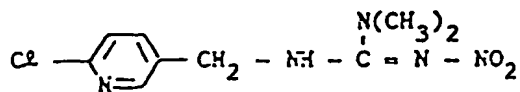
ist, worin

R⁵ und R⁶ Wasserstoff oder Methyl sind,
Z 2-Chlor-5-pyridyl ist.

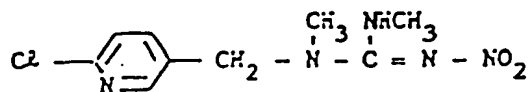
4. Verbindungen der Formel (I) nach Anspruch 1 mit den folgenden individuellen Strukturen:
3-(2-Chlor-5-pyridylmethyl)-1-methyl-2-nitroguanidin der folgenden Formel



3-(2-Chlor-5-pyridylmethyl)-1,1-dimethyl-2-nitroguanidin der folgenden Formel

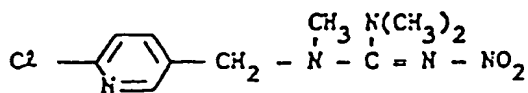


3-(2-Chlor-5-pyridylmethyl)-1,3-dimethyl-2-nitroguanidin der folgenden Formel

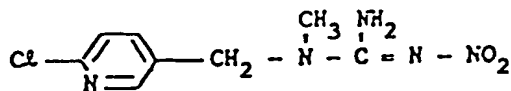


EP 0 375 907 B1

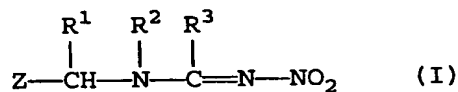
3-(2-Chlor-5-pyridylmethyl)-1,1,3-trimethyl-2-nitroguanidin der folgenden Formel



3-(2-Chlor-5-pyridylmethyl)-3-methyl-2-nitroguanidin der folgenden Formel



5. Verfahren zur Herstellung von Nitro-Verbindungen der Formel (I)



worin

R¹ und R² Wasserstoff oder C₁₋₄-Alkyl sind,
R³ -S-R⁴ oder

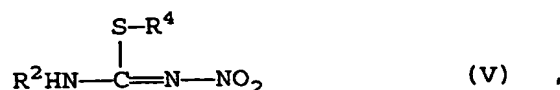


R⁴ ist, worin C₁₋₄-Alkyl ist
R⁵ und R⁶ Wasserstoff oder C₁₋₄-Alkyl sind,
Z 3-Pyridyl, 4-Pyridyl oder 5-Thiazolyl ist, das durch Halogen oder C₁₋₄-Alkyl substituiert sein kann,

mit Ausnahme von 3-(3-pyridylmethyl)-2-nitroguanidin und 3-(4-Pyridylmethyl)-2-nitroguanidin, dadurch gekennzeichnet, daß

EP 0 375 907 B1

(a) (in dem Fall, in dem R³ -S-R⁴ ist,) Verbindungen der Formel (V)



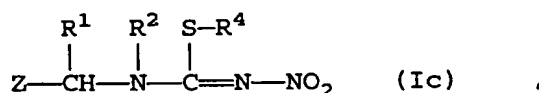
in der R² und R⁴ die oben angegebenen Bedeutungen haben, mit Verbindungen der Formel (VI)



in der R¹ und Z die oben angegebenen Bedeutungen haben und Hal ein Halogen-Atom darstellt, in Gegenwart inerter Lösungsmittel und gegebenenfalls in Gegenwart säurebindender Mittel umgesetzt werden, (b) (in dem Fall, in dem R³



ist,) Verbindungen der Formel (Ic)



worin R¹, R², R⁴ und Z die oben angegebenen Bedeutungen haben, mit Verbindungen der Formel (IV)



worin R⁵ und R⁶ die oben angegebenen Bedeutungen haben, in Gegenwart inerter Lösungsmittel umgesetzt werden.

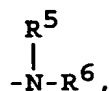
6. Insektizide Zusammensetzungen, dadurch gekennzeichnet, daß sie wenigstens eine Nitro-Verbindung der Formel (I) gemäß den Ansprüchen 1 bis 4 enthalten.
7. Verfahren zur Herstellung insektizider Zusammensetzungen, dadurch gekennzeichnet, daß Nitro-Verbindungen der Formel (I) gemäß den Ansprüchen 1 bis 4 mit Streckmitteln und/oder oberflächenaktiven Mitteln vermischt werden.
8. Verfahren zur Bekämpfung schädlicher Insekten, dadurch gekennzeichnet, daß Nitro-Verbindungen der Formel (I) gemäß den Ansprüchen 1 bis 4 auf schädliche Insekten und oder deren Lebensraum zur Einwirkung gebracht werden.

Revendications

1. Composés nitrés de formule (I)



dans laquelle R¹ et R² représentent de l'hydrogène ou un groupe alkyle en C₁ à C₄,
R³ est un groupe -S-R⁴ ou



où R⁴ est un groupe alkyle en C₁ à C₄, R⁵ et R⁶ représentent de l'hydrogène ou un groupe alkyle en C₁ à C₄.
Z est un groupe 3-pyridyle, 4-pyridyle ou 5-thiazolyle qui peut être substitué par un halogène ou par un radical alkyle en C₁ à C₄, à l'exception de la 3-(3-pyridylméthyl)-2-nitroguanidine et de la 3-(4-pyridylméthyl)-2-nitroguanidine.

2. Composés de formule (I) suivant la revendication 1, dans laquelle R¹ et R² représentent de l'hydrogène ou un groupe méthyle,
R³ est un groupe



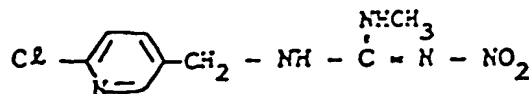
dans laquelle R⁵ et R⁶ représentent de l'hydrogène ou un groupe méthyle,
Z est un groupe 2-chloro-5-pyridyle ou 2-chloro-5-thiazolyle.

3. Composés de formule (I) suivant la revendication 1, dans laquelle R¹ et R² représentent de l'hydrogène ou un groupe méthyle,
R³ est un groupe



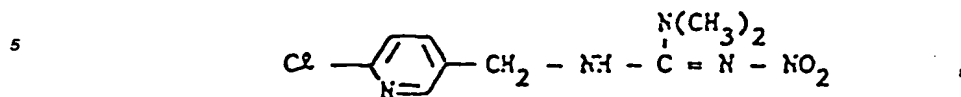
dans lequel R⁵ et R⁶ représentent de l'hydrogène ou un groupe méthyle,
Z est un groupe 2-chloro-5-pyridyle.

4. Composés de formule (I) suivant la revendication 1, présentant les structures individuelles suivantes :
3- (2-chloro-5-pyridylméthyl)-1-méthyl-2-nitroguanidine représentée par la formule suivante

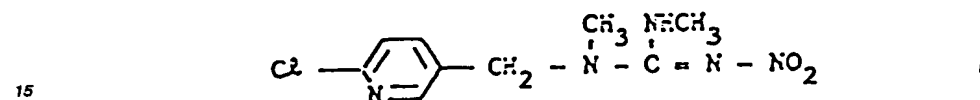


EP 0 375 907 B1

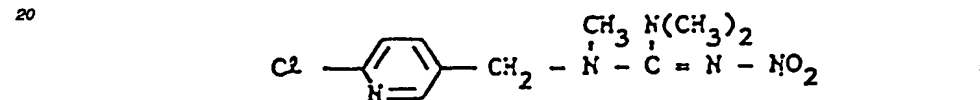
3-(2-chloro-5-pyridylméthyl)-1,1-diméthyl-2-nitroguanidine représentée par la formule suivante :



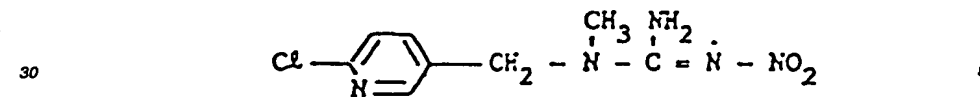
3-(2-chloro-5-pyridylméthyl)-1,3-diméthyl-2-nitroguanidine représentée par la formule suivante :



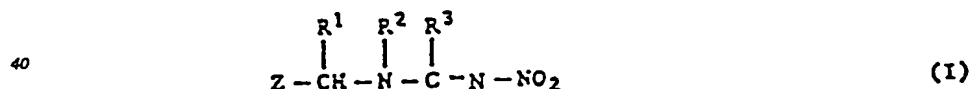
3-(2-chloro-5-pyridylméthyl)-1,1,3-triméthyl-2-nitroguanidine représentée par la formule suivante



3-(2-chloro-5-pyridylméthyl)-3-méthyl-2-nitroguanidine représentée par la formule suivante



5. Procédé de production de composés nitrés de formule (I)



dans laquelle R¹ et R² représentent de l'hydrogène ou un groupe alkyle en C₁ à C₄.

R³ est un groupe -S-R⁴ ou



où R⁴ est un radical alkyle en C₁ à C₄, R⁵ et R⁶ représentent de l'hydrogène ou un radical alkyle en C₁ à C₄.

Z est un groupe 3-pyridyle, 4-pyridyle ou 5-thiazolyle qui peut être substitué par un halogène ou un radical alkyle en C₁ à C₄.

à l'exception de la 3-(3-pyridylméthyl)-2-nitroguanidine et de la 3-(4-pyridylméthyl)-2-nitroguanidine dans lequel

EP 0 375 907 B1

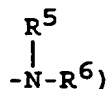
a) (au cas où R³ est un groupe -S-R⁴)
des composés de formule (V)



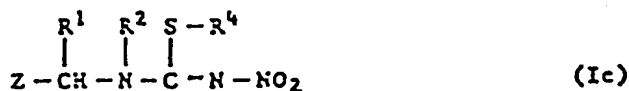
dans laquelle R² et R⁴ ont les mêmes définitions que ci-dessus,
sont amenés à réagir avec des composés de formule (VI)



dans laquelle R¹ et Z ont les mêmes définitions que ci-dessus et
Hal représente un atome d'halogène,
en présence de solvants inertes et lorsque cela convient, en présence d'accepteurs d'acides,
b) (au cas où R³ représente



des composés de formule (Ic)



dans laquelle R¹, R², R⁴ et Z ont les mêmes définitions que ci-dessus,
sont amenés à réagir avec les composés ci-dessus de formule (IV),



dans laquelle R⁵ et R⁶ ont la même définition que ci-dessus, en présence de solvants inertes.

6. Compositions insecticides, caractérisées en ce qu'elles contiennent au moins un composé nitré de formule (I) suivant les revendications 1 à 4.
7. Procédé de préparation de compositions insecticides, caractérisé en ce que des composés nitrés de formule (I) suivant les revendications 1 à 4 sont mélangés avec des diluants et/ou des agents tensio-actifs.
8. Procédé pour combattre des insectes nuisibles par application à ces insectes ou à leur habitat de composés nitrés de formule I suivant les revendications 1 à 4.